

## UNITED STATE DEPARTMENT OF COMMERCE

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Washington, D.C. 20231

APPLICATION NO. FILING DATE FIRST NAMED INVENTOR ATTORNEY DOCKET NO 09/281,474 03/30/99 RAJOPADHYE M DM-6958 **EXAMINER** HM12/1220 DAVID H. VANCE JONES, D DUPONT PHARMACEUTICALS COMPANY ART UNIT PAPER NUMBER C/O E. I. DU PONT DE NEMOURS AND CO. LEGAL - PATENTS-1007 MARKET STREET 1619 WILMINGTON DE 19898 **DATE MAILED:** 

Please find below and/or attached an Office communication concerning this application or proceeding.

**Commissioner of Patents and Trademarks** 

12/20/00

	Application No.	Applicant(s)
Office Action Summary	09/281,474	RAJOPADHYE ET AL.
	Examiner	Art Unit
	D. L. Jones	1619
The MAILING DATE of this communication appe Period for Reply	ears on the cover sheet with the c	orrespondence address
A SHORTENED STATUTORY PERIOD FOR REPL'THE MAILING DATE OF THIS COMMUNICATION.	Y IS SET TO EXPIRE <u>1</u> MONTH	(S) FROM
<ul> <li>Extensions of time may be available under the provisions of 37 after SIX (6) MONTHS from the mailing date of this commun</li> <li>If the period for reply specified above is less than thirty (30) day be considered timely.</li> <li>If NO period for reply is specified above, the maximum statutor communication.</li> <li>Failure to reply within the set or extended period for reply will, the Status</li> </ul>	ication. ys, a reply within the statutory minimum of y period will apply and will expire SIX (6)	of thirty (30) days will  MONTHS from the mailing date of this
1) Responsive to communication(s) filed on	·	
2a) This action is FINAL. 2b) Th	nis action is non-final.	
3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213.		
Disposition of Claims  4) ☐ Claim(s) 1-51 is/are pending in the application  4a) Of the above claim(s) is/are withdra  5) ☐ Claim(s) is/are allowed.  6) ☐ Claim(s) is/are rejected.  7) ☐ Claim(s) is/are objected to.	awn from consideration.	
8)⊠ Claims <u>1-51</u> are subject to restriction and/or of Application Papers	election requirement.	
9) The specification is objected to by the Examin	er.	
10) The drawing(s) filed on is/are objected		
11) ☐ The proposed drawing correction filed on is: a) ☐ approved b) ☐ disapproved.		
12) ☐ The oath or declaration is objected to by the E	xaminer.	
Priority under 35 U.S.C. § 119		
13) Acknowledgment is made of a claim for foreign	n priority under 35 U.S.C. § 119(	(a)-(d).
a) ☐ All b) ☐ Some * c) ☐ None of the CERTIF	FIED copies of the priority docum	nents have been:
2. received in Application No. (Series Cod	le / Serial Number)	
3. received in this National Stage applicati	on from the International Bureau	(PCT Rule 17.2(a)).
* See the attached detailed Office action for a list	of the certified copies not receive	red.
14) Acknowledgement is made of a claim for dom	estic priority under 35 U.S.C. & 1	119(e).
Attachment(s)		
<ul> <li>14) Notice of References Cited (PTO-892)</li> <li>15) Notice of Draftsperson's Patent Drawing Review (PTO-948)</li> <li>16) Information Disclosure Statement(s) (PTO-1449) Paper No(s)</li> </ul>	18) Notice of Inform	ary (PTO-413) Paper No(s) al Patent Application (PTO-152)

Application/Control Number: 09/281,474

Art Unit: 1619

## **RESTRICTION INTO GROUPS**

- 1. Restriction to one of the following inventions is required under 35 U.S.C. 121:
  - I. Claims 1-10, 50, and 51, drawn to compounds and methods wherein the compound comprises a chelator and peptide-targeting moiety, classified in class 424, subclass 1.69.
  - II. Claims 11-37, 48, and 49, drawn to compositions and methods wherein the compositions comprise a metal chelator and a targeting moiety, classified in class 534, subclass 7.
  - III. Claims 38-43, 46, and 47, drawn to a surfactant, and targeting moiety, classified in class 516, subclass 22.
  - Claims 44 and 45, drawn to an ultrasound agent, classified in class 424, subclass 9.5.
- 2. Inventions II III, IV, and I are unrelated. Inventions are unrelated if it can be shown that they are not disclosed as capable of use together and they have different modes of operation, different functions, or different effects (MPEP § 806.04, MPEP § 808.01). In the instant case, the different inventions while all encompassed under the very broad concept of pharmaceutical agents and methods of use thereof, the requirements for examining each group is different. In other words, four distinct groups of inventions for examining each group is different. In other words, four distinct groups of inventions for examining each group is different. In other words, four distinct groups of inventions for examining each group is different. In other words, four distinct groups of inventions for examining each group is different. In other words, four distinct groups of inventions for examining each group is different. In other words, four distinct groups of inventions for examining each group is different. In other words, four distinct groups of inventions for examining each group is different. In other words, four distinct groups of inventions for examining each group is different. In other words, four distinct groups of inventions for examining each group is different.

Art Unit: 1619

The above groups themselves are inclusive of patentably distinct subject matter. Accordingly, along with the election of one of the above groups, the following action is also taken.

The inventions are distinct from one another for the reasons set forth above. Hence, since these inventions are distinct and have acquired a separate status in the art because of their recognized divergent subject matter, restriction for examination purposes as indicated is proper.

## **ELECTION OF SPECIES**

5. Claims 1-51 are generic to a plurality of disclosed patentably distinct species comprising, for example, (1) the compounds and method of use of the compounds of Group I as set forth in independent claim 1; (2) the compositions and method of use of the compositions of Group II as set forth in independent claim 11; (3) the complexes of independent claim 38; and (4) the agents of independent claims 44 and 45.

Furthermore, a burdensome search would be necessary because of the numerous variables and substitutents present in the invention. In particular, the targeting moiety may be a peptide or peptide mimetic selected from EGFR, FGFR, PDGFR, FIk-1/KDR, FIt-1, Tek, Tie, neuropilin-1, endoglin, endosialin, and Ax1 to name a few. The variable Q may be a peptide independently selected from the group defined in claim 3 having variables L, M, K, R1, R2, R3, R4, R5, K', and/or M' or those set forth in claim 6. The

Application/Control Number: 09/281,474

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Art Unit: 1619

chelator, Ch, may be selected from those in claim 3 (see pages 178-179) containing the variables A1-A8 and E.

Applicant is required under 35 U.S.C. 121 to elect a single disclosed species, even though this requirement is traversed.

- <u>Notes</u>: (A) The Examiner respectfully requests that the Applicant assign each variable the appropriate value when electing a species (i.e., X = hydrogen; Z = nitrogen; Y = -CH3; etc.); and
- (E) The election of species has to be within the elected group as set forth above.
- 6. Should applicant traverse on the ground that the species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C. 103(a) of the other invention.
- 7. Upon the election of a single disclosed species (e.g., the first species of claim 17), a generic concept inclusive of the elected species will be identified by the Examiner for examination along with the elected species. Moreover, whatever, specific complex is ultimately elected, Applicant's are required to list all claims readable thereon.

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**Art Unit: 1619** 

- 8. A telephone call was not made to request an oral election to the restriction requirement due to the complexity of the species and restriction above.
- 9. Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).
- 10. Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a petition under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).
- 11. Any inquiry concerning this communication or earlier communications from the examiner should be directed to D. L. Jones whose telephone number is (703) 308-4640. The examiner can normally be reached on Monday Friday, 6:45 a.m. 3:15 p.m..

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Diana Dudash can be reached on (703) 308-2328. The fax phone numbers for the organization where this application or proceeding is assigned are (703) 308-4556 for regular communications and (703) 308-4556 for After Final communications.

Art Unit: 1619

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is (703) 308-1235.

D. L. Jones

Primary Examiner

Art Unit 1619

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December 12, 2000